

EXHIBIT 2

CONFIDENTIAL

Xu, Ren-He

July 3, 2019

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UNITED STATES DISTRICT COURT
DISTRICT OF MASSACHUSETTS

ASTELLAS INSTITUTE FOR)	
REGENERATIVE MEDICINE, AND)	
STEM CELL & REGENERATIVE)	
MEDICINE INTERNATIONAL, INC.,)	
)	
Plaintiffs,)	
)	
vs.)	C.A. No.
)	1:17-cv-12239
IMSTEM BIOTECHNOLOGY, INC.,)	
XIAOFANG WANG, and REN-HE XU,)	
)	
)	
Defendants.)	
_____)	

CONFIDENTIAL - PROTECTIVE ORDER

VIDEOTAPED DEPOSITION OF REN-HE XU, taken on
behalf of the Plaintiffs, at 12670 High Bluff Drive,
San Diego, California, commencing at 9:19 a.m. and
ending at 5:10 p.m., Wednesday, July 3, 2019.

Reported by:
Audrey L. Ricks
CSR No. 12098, CCR, RPR, CLR

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46 (Pages 178 to 181)

<p style="text-align: right;">178</p> <p>1 collaborative data, the data from us, especially the 2 in vivo data, in international conference. 3 Secondly, they had a plot to block our 4 publication. So our collaboration was, in my mind 5 was all about the publication in good quality 6 journal, but they had a plot to prevent that. And 7 they took -- they wanted to take all the credit. So 8 I feel unfortunate to collaborate with them. 9 BY MR. FRAZIER: 10 Q But you were given cells and a protocol; 11 correct? 12 MR. SHANNON: Objection. 13 THE WITNESS: I said I feel unfortunate. 14 If I didn't get the cells from them, I would have 15 our own TMSC method as well. 16 BY MR. FRAZIER: 17 Q But you didn't describe the TMS -- the 18 TMSC method in this grant application; right? 19 A At that time we didn't include the -- that 20 method. That was in -- under filing, I think, maybe 21 at that time. I'm not very clear. 22 First -- secondly, it's because, as I 23 said, no matter which method you make, the grant 24 application is the same. You -- you are targeting a 25 highly devastating disease in the west, and using a</p>	<p style="text-align: right;">180</p> <p>1 inhibit or inactivate the mitosis of the cells. The 2 cells still have effect. 3 This is very good news to patients. So 4 it's dramatically reduced the bio safety. You don't 5 need to concern the cells will proliferate in your 6 body. Those cells are from an ES cell line, right? 7 You don't want that. So this is great news. 8 BY MR. FRAZIER: 9 Q Other people had already 10 mitotically-inactivated MSC cells before using them 11 in animal models; correct? 12 A I don't think so. 13 Q Have you studied the literature on that? 14 A We searched around and we didn't see 15 similar reports before. 16 That's why I feel Xiaofang is a really 17 brilliant scientist. I was very fortunate to have 18 him as my postdoc, so he really had good background 19 in immunology training and also he capture a lot 20 from my stem cell research and developed very fast. 21 He's a very good scientist. 22 MR. FRAZIER: All right. Let me show you 23 what's been previously marked as Exhibit 50. 24 (Previously marked Exhibit 50 25 reviewed by the witness.)</p>
<p style="text-align: right;">179</p> <p>1 new cell type. So that's -- and especially this new 2 cell type we found it has much lower 3 pro-inflammatory factor IO6 and we could also use 4 mitotically-inactivated cells for the treatment. 5 These are all blessing, you know, for patients. So 6 that's -- those ideas matter. 7 Also you use MSC to treat the disease, 8 there's already tons of papers, tons of ideas. But 9 to use this ES-derived MSC to treat this disease 10 with low prolif -- pro-inflammatorous cytokine IO6 11 and mitotically-inactivated cells instead of 12 mitotically-active cells, this significantly reduced 13 the bio safety concern. 14 Q Right. And -- and you obviously were not 15 the first one to come up with the idea of 16 mitotically inactivating stem cells; correct? 17 A Mitotically inactivating cells is not new, 18 but mitotically inactivated ES-derived MSC is new. 19 It's important. 20 Q It was only new because they were a new 21 cell type as taught to you by ACT; right? 22 MR. SHANNON: Objection. 23 THE WITNESS: It's new because we tell -- 24 clearly tell people that mitotically inactivation 25 doesn't matter for the cell effects. So you can</p>	<p style="text-align: right;">181</p> <p>1 BY MR. FRAZIER: 2 Q My question is, do you recognize what 3 Exhibit 50 is? 4 A No. First time to me. 5 Q You haven't seen this one before? 6 A No. 7 Q Okay. All right. Are you aware that 8 there were various presentations of information made 9 to potential investors in ImStem? 10 A I knew they were looking for some more 11 investment. Yeah. 12 Q All right. But you didn't review either 13 the -- the business plan or the slides that are 14 attached to Exhibit 50? 15 A No. 16 MR. FRAZIER: Let's see. There's another 17 one in the stack there in front of you that is 18 Exhibit 66. 19 (Previously marked Exhibit 66 20 reviewed by the witness.) 21 BY MR. FRAZIER: 22 Q You see Exhibit 66 is another collection 23 of e-mails, this time from August of 2013? 24 A Yeah. There's -- it's talking about mice. 25 Q Yeah. So you see it begins on page 31134</p>